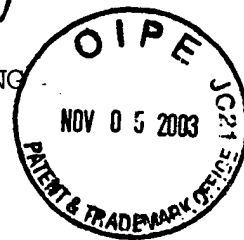


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ESSENTIAL GENES FROM C. ALBICANS AND A METHOD FOR SCREENING
ANTIMYCOTIC SUBSTANCES USING SAID GENES



Field of the Invention

5 The present invention relates to a method for
screening for antimycotic substances in which essential
genes from mycetes, particularly from Candida albicans (C.
albicans) as well as functionally similar genes from other
pathogenic mycetes, or the corresponding encoded proteins,
10 are used as targets. The invention also relates to specific
C. albicans genes.

Background of the Invention

 The spectrum of known fungal infections stretches from
15 fungal attack of skin or nails to potentially hazardous
mycotic infections of the inner organs; Such infections and
resulting diseases are known as mycosis. Antimycotic
substances (fungistatic or fungicidal) are used for
treatment of mycosis. However, up to now, relatively few
20 substances with pharmacological effects are known, such as
Amphotericin B, Nystatin, Pimaricin, Griseofulvin,
Clotrimazole, 5-fluoro-cytosine and Batraphene. The drug
treatment of fungal infections is extremely difficult, in
particular because both the host cells and the mycetes, are
25 eucaryotic cells. Administration of drugs based on known
antimycotic substances results therefore often in undesired
side-effects, for example Amphotericin B has a nephrotoxic
effect. Therefore, there is a strong need for
pharmacologically efficient substances usable for the
30 preparation of drugs, which are suitable for prophylactic
treatments of immunodepressive states or for the treatment
of an existing fungal infection. Furthermore, the
substances should exhibit a specific spectrum of action in
order to selectively inhibit the growth and proliferation
35 of mycetes without affecting the treated host organism.

Summary of the Invention

The aim of the present invention is to provide a method for the identification of antimycotic substances and especially for the identification of anti-Candida substances. An essential feature of this method is that essential genes from mycetes are used as targets for the screening.

The present invention thus concerns a method for screening antimycotic substances wherein an essential gene from mycetes or a functionally similar gene in another pathogenic mycete, or the corresponding encoded protein, is used as target and wherein the essential gene is selected from the group consisting of CaOR110, CaMR212, CaNL256, CaBR102, CaIR012, CaDR325 and CaJL039.

Brief Description of the Figures

Figure 1 shows the comparison of the deduced protein of YNL256 gene with YNL256.

Figure 2 shows the comparison of the deduced YBR102 gene with YNL102.

Figure 3 shows the alignment of ScOR110-homologous fragment with Candida albicans CaOR110 sequence.

Figure 4 is a map of CaOR110.

Figure 5 is a map of the CaOR110 variant.

Figures 6A, 6B and 6C show the alignment of the original CaOR110 and splice variant.

Figure 7 shows a map of CaDR325.

Detailed Description of the Invention

According to one embodiment of the method of the invention mycete cells which express the essential gene, or a functionally similar mycete gene, to a different level are incubated with the substance to be tested and the growth inhibiting effect of the substance is determined.

According to another embodiment, said target gene or the corresponding target gene encoded protein is contacted

According to another embodiment, said target gene or the corresponding target gene encoded protein is contacted in vitro with the substance to be tested and the effect of the substance on the target is determined.

5 According to another embodiment, the screened substances inhibit partially or totally the functional expression of the essential genes or the functional activity of the encoded proteins.

According to one embodiment the screened substances
10 partially or totally inhibit the activity of dihydropteroate synthase (DHPS) and/or 7,8-dihydro-6-hydroxymethylpterin-pyrophosphokinase (HPPK).

According to another embodiment, the mycete species are selected from the group comprising Basidiomycetes,
15 Ascomycetes and Hyphomycetes.

According to another embodiment of the method of the invention said functional similar genes are essential genes from *Candida* Spp., preferably *Candida albicans*, or from *Aspergillus* Spp., preferably from *Aspergillus fumigatus*.

20 According to another aspect, the present invention concerns a polynucleotide having the sequence as depicted in SEQ ID No. 2, SEQ ID No. 5, SEQ ID No. 8, SEQ ID No. 10, SEQ ID No. 13, SEQ ID No. 15, SEQ ID No. 17 or SEQ ID No. 20, preferably SEQ ID No. 2, SEQ ID No. 5, SEQ ID No. 8,
25 SEQ ID No. 13, SEQ ID No. 14 or SEQ ID No. 17, homologs thereof and functional fragments thereof.

According to another aspect, the present invention concerns a gene which is CaOR110, CaMR212, CaNL256, CaBR102, CaIR012, CaDR325 or CaJL039, preferably CaOR110,
30 CaMR212, CaNL256, CaBR102 or CaIR012, or a functionally similar gene or a functional fragment thereof.

According to this embodiment, the functionally similar gene or homologous polynucleotide has a sequence identity, at the nucleotide level, with CaOR110, CaMR212, CaNL256,
35 CaBR102, CaIR012, CaDR325 or CaJL039, respectively, of at least 50%, preferably of at least 60%, and most preferably

of at least 70%. A functional fragment is a polynucleotide fragment that will retain the functionality of the starting product (nucleotide or gene). One example is the CaOR110 splice variant (which is also homologous to the original gene, with about 90% identity).

According to another embodiment, the functionally similar gene has a sequence identity, at the amino-acid level, with CaOR110, CaMR212, CaNL256, CaBR102, CaIR012, CaDR325 or CaJL039, respectively, encoded protein (s) of at least 40%, preferably of at least 50%, more preferably of at least 60% and most preferably of at least 70%.

These figures given for the gene apply mutatis mutandis to the polynucleotide, as far as homology and similarity.

According to another aspect, the present invention covers the protein(s) encoded by CaOR110, CaMR212, CaNL256, CaBR102, CaIR012, CaDR325 or CaJL039, respectively, gene(s) or by a functionally similar gene, or a functional polypeptidic fragment thereof.

According to another aspect, the present invention provides a plasmid containing CaOR110, CaMR212, CaNL256, CaBR102, CaIR012, CaDR325 or CaJL039, respectively, gene (s), a functionally similar gene or a functional fragment thereof. According to another aspect, the present invention provides a plasmid (bacteria containing same) deposited at the CNCM (Institut Pasteur, Paris) on 98/08/13, with the accession numbers 1-2065, I-2063 and I-2064, corresponding to the CaNL256, CaBR102 and CaIR012 genes, respectively.

According to another aspect, the present invention provides a plasmid (bacteria containing same) deposited at the DSMZ (Deutsche Sammlung von Mikroorganismen und Zellkulturen, Germany) on 99/08/06 with the accession numbers DSM 12977, DSM 12976, DSM 12978 and DSM 12979, corresponding to the CaDR325, CaOR110, CaOR110 splice variant and CaMR212, respectively.

According to another aspect, the present invention provides a kit for diagnosis of fungal infections comprising a gene selected from the group consisting of CaOR110, CaMR212, CaNL256, CaBR102, CaIR012, CaDR325 and CaJL039, a functionally similar gene thereof, a functional fragment thereof, the corresponding encoded protein or a functional polypeptide fragment thereof.

According to another aspect, the present invention provides an antibody directed against the protein encoded by the CaOR110, CaMR212, CaNL256, CaBR102, CaIR012, CaDR325 or CaJL039, respectively, gene(s) or by a functionally similar gene, or a polypeptide fragment thereof.

According to another aspect, the present invention provides a polynucleotide obtainable by the process comprising the following steps:

- (i) selecting an essential gene from *Saccharomyces cerevisiae*;
 - (ii) comparing the sequence of said gene with *Candida Albicans* genome sequences;
 - (iii) deducing homologous oligonucleotides regions ;
 - (iv) PCR amplifying the thus-obtained oligonucleotides;
 - (v) using the amplimers of step (iv) for detecting the complete gene of interest:
- the amplimers of step (iv) are used as a probe for detecting the complete gene of interest from a *Candida albicans* genomic or cDNA library; or
- the complete gene is obtained by 3' and 5' extension of the amplimer, e. g. by using a PCR method.

According to the invention, the first step is to identify said essential genes and starting from these thus identified genes, essential genes from other pathogenic mycetes can be identified. For practical purposes, essential genes from *S. cerevisiae* are first identified and

starting from them, essential genes from other pathogenic fungus, especially from *Candida*, are obtained.

The present invention thus discloses the identification of essential genes from *C. albicans* and
5 their use in a method for the screening of antimycotic substances, especially anti-*Candida* substances.

In order to identify essential genes of *S. cerevisiae*, individual genomic genes are eliminated through homologous recombination. If the DNA segment thus eliminated concerns
10 an essential gene, then the deletion is lethal for the *S. cerevisiae* cells in haploid form.

A method, wherein the studied *S. cerevisiae* gene is replaced by a marker gene can be used to generate the corresponding genomic deletion of *S. cerevisiae* and to
15 determine the *S. cerevisiae* cells containing the deletion.

As a selection marker a dominant selection marker (e. g. kanamycin resistance gene) or an auxotrophic marker can for example be used. As an auxotrophic marker, it is possible to use genes coding for key enzymes of amino acid
20 or nucleic base synthesis. For example, one can use as a selection marker the following genes from *S. cerevisiae* : gene encoding for the metabolic pathway of leucine (e. g. LEU2-gene), histidine (e. g. HIS3-gene) or tryptophan (e. g. TRP-1 gene) or for the nucleic base metabolism of uracil
25 (e. g. URA3-gene).

Auxotrophic *S. cerevisiae* strains can be used. These auxotrophic strains can only grow on nutritive media containing the corresponding amino acids or nucleotide bases. All laboratory *S. cerevisiae* strains, containing
30 auxotrophic markers can for instance be used. When diploid *S. cerevisiae* strains are used, then the corresponding marker gene must be homozygously mutated. Strain CEN. PK2 or isogenic derivatives thereof can be used.

Strains containing no suitable auxotrophic marker can
35 also be used such as prototrophic *S. cerevisiae* strains. Then a dominant selection marker e. g. resistance

gene, such as kanamycin resistance gene can be used. A loxP-KanMX-loxP cassette can advantageously be used for this purpose.

For the homologous recombination replacing the whole
 5 DNA sequence or part thereof of a *S. cerevisiae* gene, DNA fragments are used wherein the marker gene is flanked at the 5'-and 3'-ends by sequences which are homologous to the 5'-and 3'-ends of the studied *S. cerevisiae* gene.

Different processes can be used for the preparation of
 10 the corresponding DNA fragments which are also appropriate for the deletion of any specific *S. cerevisiae* gene. A linear DNA-fragment is used for the transformation of the suitable *S. cerevisiae* strain. This fragment is integrated into the *S. cerevisiae* genome by homologous recombination.
 15 These processes include:

1. "Conventional method" for the preparation of deletion cassettes (Rothstein, R. J. (1983) *Methods in Enzymology*, Vol. 101, 202-211).

2. "Conventional Method" using the PCR technique
 20 ("modified conventional method").

3. SFH (short flanking homology)-PCR method (Wach, A. et al. (1994) *Yeast* 10: 1793-1808; Gültner, U. et al. (1996) *Nucleic Acids Research* 24: 2519-2524).

1. In the "conventional method" for the preparation of
 25 deletion cassettes in the *S. cerevisiae* genome, the gene to be studied is either already present in an appropriate vector or is integrated in such a vector. With this method, any pBR-pUC-and [pBluescript®-derivatives] can be used for example. A major part of the target gene sequence is
 30 eliminated from the vector, for instance using appropriate restriction sites, conserving however the 3'-and 5'-regions of the studied gene inside the vector. The selected marker gene is integrated between the remaining regions.

2. In the modified form of this "conventional method",
 35 PCR is used. This method allows amplification of the 3'-and 5'-terminal regions of the coding sequence of the

studied *S. cerevisiae* gene. This method amplifies selectively both terminal regions of the studied gene, therefore, two PCR-reactions must be carried out for each studied gene, amplifying once the 5'-end, and once the 3'-end of the gene. The length of the amplified terminal DNA-fragment depends on the existing restriction sites. The amplified terminal ends of the studied gene have generally a length of 50 to 5000 base pairs (bp), preferably a length between 500 and 1000 bp.

As template for the PCR-reactions, genomic DNA of *S. cerevisiae* or wild-type genes can be used. The primer-pairs (a sense and an antisense primer, respectively) are constructed so that they correspond to the 3'-end and the 5'-end sequence of the studied *S. cerevisiae* gene.

Especially, the primer is selected such as to allow its integration by way of appropriate restriction sites.

As vectors, pBR-pUC-and pBluescript®-derivatives can be used. In particular vectors already containing a gene encoding the selection marker, are appropriate. In

particular, vectors can be used, which contain genes of the selection marker HIS3, LEU2, TRP1 or URA3.

The DNA segments of the studied *S. cerevisiae* gene, obtained by PCR, are integrated in the vector at both sides of the selection marker, so that subsequently, as in the "conventional method", the selection marker is flanked on both ends by DNA sequences which are homologous to the studied gene.

3. Homologous recombination in *S. cerevisiae* takes place in a very efficient and precise manner and the length of the DNA sequence homologous to the studied *S. cerevisiae* gene flanking the selection marker gene can in fact be considerably shorter than with the "modified conventional method". The flanking ends homologous to the studied *S. cerevisiae* gene need to present a length of only about 20-60 bp, preferably 30-45 bp. The SFH-PCR method is

particularly advantageous as the laborious cloning step can be obviated.

A PCR reaction is carried out on a DNA-template containing the gene for the selection marker to be used, wherein the primers are constructed so that the DNA sequence of the sense primer is homologous to the 5'-end of the selection marker sequence and so that the primer presents in addition at its 5'-end a region of preferably 40 nucleotides, which corresponds to the 5'-terminal sequence of the studied *S. cerevisiae* gene. The antisense primer is constructed in an analogous manner, i. e. it is complementary to the 3'-end of the gene sequence of the selection marker, wherein this primer contains at its 5'-end a region of also preferably 40 nucleotides, which corresponds to the complementary strand of the 3'-terminal coding sequence of the studied gene.

For the amplification of *S. cerevisiae* genes to be studied by the SFH-PCR method, vectors containing the gene for the auxotrophic marker or selection marker can be used. Especially, plasmid pUG6 is used as the template. This plasmid contains a loxP-KanMX-loxP cassette (Gültner, U. et al. (1996) Nucleic Acids Research 24: 2519-2524). In other terms, the Kanamycin resistance gene is flanked at both ends by a loxP sequence (loxP-KanMX-loxP cassette). This cassette is advantageous in that the Kanamycin resistance gene can be eventually eliminated from the *S. cerevisiae* genome after integration of the loxP-KanMX-loxP cassette into the *S. cerevisiae* gene to be studied. Cre-recombinase of bacteriophage P1 can be used for this purpose. Cre-recombinase recognizes the loxP sequences and induces elimination of the DNA located between the two loxP sequences by a homologous recombination process. As a result only one loxP sequence remains and the so-called marker regeneration occurs, i. e. the *S. cerevisiae* strain may be transformed again using the loxP-KanMX-loxP cassette. This is particularly advantageous, when at least

two functionally similar genes are to be deleted in order to obtain a lethal phenotype.

With the PCR-method, the PCR reaction primers are at the 3'-end a preferably 20 nucleotide long sequence, which is homologous to the sequence situated left and/or right of the loxP-KanMX-loxP cassette, and at the 5'-end a preferably 40 nucleotide long sequence, which is homologous to the terminal ends of the gene to be studied.

Using the three methods, one obtains linear deletion cassettes containing the gene encoding the selection marker, which is flanked on both sides by homologous sequences of the gene to be studied. The deletion cassettes are used for the transformation of diploid *S. cerevisiae* strains. The diploid strain *S. cerevisiae* CEN. PK2 (Scientific Research & Development GmbH, Oberursel) can be used for example for this purpose.

[CEN. PK2 Mata/MAT a ura3-52/ura3-52 leu2-3,112/leu2-3,112his3A1/his3A1 trp1-289/trp1-289 MAL2-8C/MAL2-8c SUC2/SUC2]

The strain CEN. PK2 is prepared and cultivated using known methods (Gietz, R. D. et al. (1992) *Nucleic Acids Research* 8: 1425; Guldener, U. et al. (1996) *Nucleic Acids Research* 24: 2519-2524).

The cells of the *S. cerevisiae* strain used are transformed according to known processes with an appropriate DNA quantity of the linear deletion cassette (e. g. Sambrook et al. 1989). Thereafter, the medium in which the cells are cultivated is replaced by a new medium, a so-called selective medium, which does not contain the corresponding amino acid (e.g. histidine, leucine or tryptophan) or nucleic base (e.g. uracil) or, when using a deletion cassette containing the kanamycin resistance gene, by a medium containing geneticin (G4180) (e. g. a complete medium (YEPD) containing geneticin). Alternatively, the transformed cells may be plated on agar plates prepared using the corresponding media. Thereby, one selects the

transformed cells, in which a homologous recombination occurred, since only those cells can grow under these modified conditions.

However, in most cases, only one of the two copies of the gene in the double chromosome set of a diploid *S. cerevisiae* strain is replaced by the DNA of the deletion cassette during the transformation, resulting in a heterozygote-diploid *S. cerevisiae* mutant strain, wherein one copy of the gene studied is replaced by a selection marker, while the other copy of the gene is maintained in the genome. This presents the advantage that in case of a deletion of an essential gene, due to the existence of the second copy of the essential gene, the mutant *S. cerevisiae* strain is still viable.

The proper integration of the deletion cassette DNA at the predetermined chromosomal gene locus (gene locus of the gene to be studied) may be checked by Southern-Blot Analysis (Southern, E. M. (1975) *J. Mol. Biol.* 98: 503-517) or by diagnostic PCR analysis using specific primers (Guldener, U. et al. (1996) *Nucleic Acids Research* 24: 2519- 2524)

The genetic separation of individual diploid cells may be monitored by tetrad analysis. To this end, reduction division (meiosis) is induced in the diploid cells, especially heterozygote mutant strains, using known methods such as nitrogen impoverishment on potassium acetate plates (Sherman, F. et al. (1986) Cold Spring Harbor Laboratory Press, Cold Spring Harbor, N. Y.; Guthrie, C. and Fink, G. R. (1991) *Methods in Enzymology*, Vol 194. Academic Press, San Diego, 3-21; Ausubel, F. M. et al. (1987) *Current Protocol in Molecular Biology* John Wiley and Sons, Inc., Chapter 13). Meiosis results only in asci with four ascospores (segregated), which can be individualized after partial enzymatic digestion of the ascospore wall with zymolyase (Ausubel et al. (1987)) by way of micromanipulators (e. g. SINGER). For example when a tetrad

analysis is carried out on a heterozygote-diploid mutant strain in which an essential gene present in the double chromosome set is replaced on one chromosome by homologous recombination, then only two segregated ascospores are viable, namely those which carry the essential gene. The two remaining segregated ascospores are not viable because they lack the essential gene.

In order to check if the genes studied by this method are really essential or if the homologous recombination leads to an alteration of an essential gene adjacent to the gene locus of the gene studied, the heterozygote diploid *S. cerevisiae* mutant strain is transformed with a centromere plasmid containing said studied gene.

A tetrad analysis is carried out on the transformants. When four instead of two viable segregates are obtained, then the studied gene contained in the centromere plasmid can complement the defect of the two non-viable haploid *S. cerevisiae* cells/mutant strains, which demonstrates that the studied *S. cerevisiae* gene is essential.

Preferably, plasmids present in low copy number, e. g. one or two copies per cell are used as centromere plasmids. For example plasmids pRS313, pRS314, pRS315 and pRS316 (Sijkorski, R. S. and Hieter, P. (1989) Genetics 122: 19- 27) or similar plasmids can be used for this purpose. Preferably, the studied genes are integrated in said plasmids including their 3'-and 5'-end non-coding regions.

Individual *S. cerevisiae* genes may be studied using the above-described method, their sequences being totally or partially known. The complete genomic sequence of *S. cerevisiae* was made accessible to the public via the WWW (World Wide Web) on April 24, 1996.

Different possibilities exist to have access to the *S. cerevisiae* genomic DNA sequence via the WWW.

MIPS (Munich information Centre of Protein Sequence) Address <http://speedY.mips.biochem.mpa.de/mips/yeast/>

SGD (Saccharomyces Genome Database, Stanford)
 Address <http://genome-www.stanford.edu/Saccharomyces> YPD (Yeast Protein Database, Cold
 Spring Harbor) Address <http://www.proteome.com/YPDhome.html>

The complete *S. cerevisiae* DNA sequence is also
 accessible via FTP (file transfer protocol) in Europe (e.
 g. at the address: [ftp.mips.embl.net](ftp:mips.embl.net)) in the U.S.A.
 (address: [genome-www.stanford.edu](ftp:genome-www.stanford.edu)) or in Japan (address:
 10 [ftp.nig.ac.jp](ftp:nig.ac.jp)).

7 essential genomic *S. cerevisiae* genes have been
 identified by this way: YDR325w, YJL039c, YOR110w, YNL256w,
 YBR102c, YIR012w and YMR212c

The essential genes of *S. cerevisiae* are then used to
 15 identify corresponding functionally similar genes in other
 mycetes.

By functionally similar genes in other mycete species,
 is meant genes which have a function similar or identical
 to that of the identified essential genes of *S. cerevisiae*.
 20 Functionally similar genes in other mycetes may, but need
 not be homologous in sequence to the corresponding
 essential *S. cerevisiae* genes. Functionally similar genes
 in other mycetes may exhibit only moderate sequence
 homology at the nucleotide level to the corresponding
 25 essential *S. cerevisiae* genes. By moderate sequence
 homology it is meant in the present invention genes having
 a sequence identity, at the nucleotide level, of at least
 50%, more preferably of at least 60% and most preferably of
 at least 70%.

30 In addition, functionally similar genes in other
 mycetes may, but need not encode proteins homologous in
 sequence to the proteins encoded by the essential *S.*
cerevisiae genes. Functionally similar proteins in other
 mycetes may exhibit moderate protein sequence homology to
 35 the proteins encoded by the essential *S. cerevisiae* genes.

By moderate protein sequence homology is meant in the present invention proteins having a sequence identity, at the amino-acid level, of at least 40%, preferably of at least 50%, more preferably of at least 60% and most preferably of at least 70%.

Genes homologous in sequence may be isolated according to known methods, for example via homologous screening (Sambrook, J. et al. (1989) Molecular Cloning Cold Spring Harbor Laboratory Press, N. Y.) or via the PCR technique using specific primers from genomic libraries and/or cDNA libraries of the corresponding mycetes.

According to one embodiment, genes homologous in sequences are isolated from cDNA libraries. In order to find out functionally similar genes in other mycetes, mRNA is isolated from mycete species to be studied according to known methods (Sambrook et al. 1989) and cDNA is synthesized according to known methods (Sambrook et al. 1989; or cDNA synthesis kits, e. g. from STRATAGENE).

The prepared cDNA is directionally integrated in a suitable expression vector.

For example, synthesis of the first cDNA strand may be carried out in the presence of primers having appropriate restriction sites in order to allow a subsequent cloning in the proper orientation with respect to the expression vector promoter. As restriction sites, any known restriction site may be used. As a primer, for instance the following primer, 50 nucleotides long may be used: 5'GAGAGAGAGAGAGAGAGAGAGAACTAGTXXXXXTTTTTTTTTTTTTTTTTT-3' (SEQ ID NO: 20)

The sequence (X) 6 represents an appropriate restriction site, for example for XhoI.

After two-strand synthesis, the cohesive ends of the double stranded cDNA are filled (blunt end) and the cDNA ends are then ligated using a suitable DNA adaptor sequence. The DNA adaptor sequence should contain a

restriction site which should be different from the restriction site used in the primer for the synthesis of the first cDNA strand. The DNA adaptor may comprise for example complementary 9-or 13-mer oligonucleotides, whose
 5 ends represent the cohesive end of a restriction site.

These ends may be for example a EcoRI-site:

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5' XXXXXGGCAGCAG 3'
3'       XCCGTGCTC 5'

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The single-stranded X in the adaptor sequence
 10 represent the cohesive end of a restriction site.

The cDNA provided with corresponding adaptor sequences is then cleaved using restriction endonuclease, whose recognition site was used in the primer for the synthesis of the first cDNA strand, for example XhoI. The cDNA thus
 15 obtained would have according to this example 3'-XhoI and 5'-EcoRI protruding ends and could be therefore directionally integrated into an expression vector cleaved with XhoI and EcoRI.

As expression vectors, among others, *E. coli*/S.
 20 *cerevisiae* shuttle vectors, i. e. vectors usable in *E. coli* as well as in *S. cerevisiae* are suitable. Such vectors may then be amplified for instance in *E. coli*. As expression vectors, those which are present in a high copy number as well as those present in a low copy number in *S. cerevisiae*
 25 cells can be used. For this purpose, for example vectors selected in the group consisting of pRS423 -pRS426 (pRS423, pRS424, pRS425, pRS426) and/or pRS313- pRS316 (pRS313, pRS314, pRS315, pRS316) (Sikorki, R. S. and Hieter, P. (1989) Genetics 122: 19-27; Christianson T. W. et al.
 30 (1992) Gene 110: 119-122) are suitable.

Expression vectors should contain appropriate *S. cerevisiae* promoters and terminators. In case they do not have these elements, the corresponding promoters and terminators are inserted in such a way that a subsequent
 35 incorporation of the generated cDNA remains possible.

Particularly suitable are the promoters of *S. cerevisiae* genes MET25, PGK1, TPI1, TDH3, ADHI, URA3. One may use promoters of the wild-type gene in non modified form as well as promoters which were modified in such a way that certain activator sequences and/or repressor sequences were eliminated. As terminators, for example the terminators of the *S. cerevisiae* genes MET25, PGK1, TPI1, TDH3, ADHI, URA3 are suitable.

According to another embodiment, genes homologous in sequence are isolated from genomic libraries. Genomic DNA libraries from mycetes can be prepared according to procedures known (for example as described in Current Protocols in Molecular Biology, John Wiley and Sons, Inc). For example, genomic DNA from mycetes can be prepared using known methods for yeast cell lysis and isolation of genomic DNA (for example commercially available kits from Biol01, Inc). The genomic DNA can be partially digested using a restriction enzyme such as Sau3AI and the fragments are size-selected by agarose gel electrophoresis. DNA fragments having for example a size of 5-10kb are then purified by classical methods (as for example, using Gene Clean kit from Biol01) and inserted in a *E. coli*/yeast shuttle vector such as YEP24 (described e. g. by Sanglard D., Kuchler K., Ischer F., Pagani J-L., Monod M. and Bille J., Antimicrobial Agents and Chemotherapy, (1995) Vol. 39 Noll, P2378-2386) cut by a restriction enzyme giving compatible ends (for example BamHI for Sau3AI-cut genomic DNA). The resulting expression library can be amplified in *E. coli*. However any known method, appropriate for the preparation of a genomic library, can be used in the present invention.

In order to find the genes in the studied mycete species, which are functionally similar to essential genes of *S. cerevisiae*, one *S. cerevisiae* essential gene is placed under control of a regulated promoter, either as an integrative (1) or extrachromosomal (2) gene.

1. For the integration of a regulated promoter in the *S. cerevisiae* genome, one replaces the native promoter of the selected essential gene by the regulated promoter, for example by homologous recombination via PCR (Guldener et al. (1996)). The homologous recombination via PCR can be carried out for example in the diploid *S. cerevisiae* strain CEN. PK2. The successful integration into one chromosome can be checked in haploid cells following tetrad analysis.

Using the tetrad analysis, one obtains four viable ascospores, wherein in two haploid segregates, the selected essential gene is placed under the control of the native promoter, while the essential gene in the two remaining segregates is placed under the control of the regulated promoter.

The last mentioned haploid segregates are used for the transformation with the cDNA or the genomic DNA present in the recombinant vector.

2. Using the extrachromosomal variant, the selected essential *S. cerevisiae* gene is first inserted in a suitable expression vector, for example a *E. coli*/ *S. cerevisiae* shuttle vector. For this purpose, the essential gene may be amplified via PCR from genomic *S. cerevisiae* DNA starting from the ATG initiation codon up to and including the termination codon. The primers used for this purpose may be constructed in such a way that they contain recognition sites for appropriate restriction enzymes, facilitating a subsequent insertion under control of a regulated promoter in an expression vector.

The recombinant expression vector with the plasmid copy of the essential *S. cerevisiae* gene under the control of a regulated promoter is subsequently used for the transcomplementation of the corresponding mutant allele. The corresponding mutant allele may be selected from the heterozygote-diploid mutant strains prepared by eliminating, partially or totally, by homologous

recombination an essential mycete gene listed above and as described above.

The expression vector with the selected essential *S. cerevisiae* gene is transformed in the corresponding
 5 heterozygote-diploid mutant strain carrying instead of the selected essential *S. cerevisiae* gene, a selection marker gene. The transformants are isolated by selection based on the auxotrophic marker contained in the expression vector used. The thus transformed heterozygote-diploid mutant
 10 strains are submitted to a tetrad analysis. One obtains four viable segregates. By retracing the corresponding markers of the mutant allele and the expression vector, the transformed wild-type segregates may be distinguished from segregates which do not contain the genomic copy of the
 15 essential gene. Segregates, which do not contain the genomic copy of the selected essential gene, are designated as trans-complemented haploid mutant strains. They are subsequently used for transformation with cDNA or genomic DNA libraries from other mycete species present in
 20 appropriate vectors.

As regulated promoters, inducible or repressible promoters may be used. These promoters can consist of naturally and/or artificially disposed promoter sequences.

As regulated promoters, for example the promoters of
 25 *GAL1* gene and the corresponding promoter derivatives, such as for example promoters, whose different UAS (upstream activation sequence) elements have been eliminated (GALS, GALL; Mumberg, J. et al. (1994) *Nucleic Acids Research* 22: 5767-5768) may be used. As regulated promoters, promoters
 30 of gluconeogenic genes may also be used, such as e. g. *FBP1*, *PCK1*, *ICL1* or parts therefrom, such as e. g. their activation sequence (UAS1 and/or UAS2) or repression sequence (URS, upstream repression sequence) (Niederacher et al. (1992), *Curr. Genet.* 22: 636-670; Proft et al. (1995)
 35 *Mol. Gen. Gent.* 246: 367-373; Schüller et al. (1992) *EMBO J.* 11: 107-114; Guarente et al. (1984) *Cell* 36: 503- 511).

A *S. cerevisiae* mutant strain modified in this manner can be cultivated under growth conditions, in which the regulated promoter is active, so that the essential *S. cerevisiae* gene is expressed. The *S. cerevisiae* cells are
5 then transformed with a representative quantity of the library containing the studied mycete species cDNA or genomic DNA. Transformants express additionally the protein whose coding sequence is present in the recombinant vector.

The method contemplates that the growth conditions may
10 be modified in such a way as to inhibit the regulated promoter, under the control of which is the selected essential gene. Especially, growth conditions may be changed by replacing the growth medium. When for example the GAL1 promoter or a derivate thereof is used, one can
15 replace the galactose-containing medium (induced state) by a glucose-containing medium (repressed state).

These modified conditions are lethal for the *S. cerevisiae* cells in which the recombinant vector does not carry the functionally similar genomic DNA or cDNA of the
20 studied mycete species. On the contrary, the *S. cerevisiae* cells in which the recombinant vector expresses a functionally similar coding sequence of the studied mycete species, are viable, since in these cells the lethal metabolic defect is complemented by the protein encoded by
25 the functionally similar gene.

The method contemplates that the recombinant vector (the plasmid) is isolated from the surviving transformants using known method (Strathern, J. N. and Higgins, D. R. (1991). Plasmids are recovered from yeast into
30 *Escherichia coli* shuttle vectors in: Guthrie, C. and Fink, G. R. Methods in Enzymology, Volume 194. Guide to yeast genetic and molecular Biology. Academic Press, San Diego, 319-329) and the cDNA or genomic DNA is analyzed using DNA-analysis methods such as DNA sequencing. (Sanger et al.
35 (1977), Proc. Natl. Acad. Sci. USA 74: 5463-5467)

Essential *S. cerevisiae* genes may thus be used for the identification of functionally similar genes and/or genes homologous in sequence in other mycetes, especially essential genes functionally similar and/or homologous in sequence in mycetes pathogenic to human, animal and plants. For this purpose for example mycetes of the classes phycomycetes or eumycetes may be used, in particular the subclasses basidiomycetes, ascomycetes, especially mehiascomycetales (yeast) and plectascales (mould fungus) and gymnascales (skin and hair fungus) or of the class of hyphomycetes, in particular the subclasses conidiosporales (skin fungus) and thallosporales (budding or gemmiparous fungus), among which particularly the species *mucor*, *rhizopus*, *coccidioides*, *paracoccidioides* (*blastomyces brasiliensis*), *endomyces* (*blastomyces*), *aspergillus*, *penicilium* (*scopulariopsis*), *trichophyton* (*ctenomyces*), *epidermophyton*, *microsporon*, *pieiraia*, *hormodendron*, *phialophora*, *sporotrichon*, *cryptococcus*, *candida*, *geotrichum* and *trichosporon*.

Of particular interest is the use of *Candida* Spp. especially *Candida albicans*, *Candida glabrata*, *Aspergillus* Spp., especially *Aspergillus fumigatus*, *Coccidioides immitis*, *Cryptococcus neoformans*, *Histoplasma capsulatum*, *Blastomyces dermatitidis*, *Paracoccidioides brasiliensis* and *Sporothrix schenckii*.

Starting from the genes of *S. cerevisiae*, identified according to the above-described method, Applicants cloned corresponding essential genes from *C. albicans* i. e. CaOR110, CaMR212, CaNL256, CaBR102, CaIR012, CaDR325 or CaJL039, by the following method.

First, oligonucleotide (s) is (are) selected in the sequence of the *S. cerevisiae* gene or a homologous *C. albicans* sequence in order to amplify the corresponding fragment of *C. albicans*. After cloning, the obtained fragment (exhibiting a sequence of about several hundred bp) is used as a probe for screening a *C. albicans*

(genomic) DNA library. The screening may include the following steps: clones were spread on dishes, covered with filters where the DNA was crosslinked to the filters, filters are hybridized, the positive colonies are then
5 detected. The selected clone (s) is (are) then sequenced.

The method contemplates that essential mycete genes are used to identify substances which may inhibit partially or totally the functional expression of these essential genes and/or the functional activity of the encoded
10 proteins. Substances may be identified in this fashion, which inhibit mycetes growth and which can be used as antimycotics, for example in the preparation of drugs.

The present invention especially covers a method for screening such inhibiting substances wherein an essential
15 gene from *C. albicans* selected from CaOR110, CaMR212, CaNL256, CaBR102, CaIR012, CaDR325 or CaJL039, or a functionally similar gene in another pathogenic mycete or the corresponding encoded protein is used as target.

By functionally similar genes in other pathogenic
20 mycete species, is meant genes which have a function similar or identical to that of the identified essential genes of *C. albicans*. Functionally similar genes in other pathogenic mycetes may, but need not be homologous in sequence to the corresponding essential *C. albicans* genes.
25 Functionally similar genes in other pathogenic mycetes may exhibit only moderate sequence homology at the nucleotide level to the corresponding essential *C. albicans* genes. By moderate sequence homology it is meant in the present invention genes having a sequence identity, at the
30 nucleotide level, of at least 50%, more preferably of at least 60% and most preferably of at least 70%.

In addition, functionally similar genes in other pathogenic mycetes may, but need not encode proteins homologous in sequence to the proteins encoded by the
35 essential *C. albicans* genes. Functionally similar proteins in other mycetes may exhibit moderate protein sequence

homology to the proteins encoded by the essential *C. albicans* genes.

By moderate protein sequence homology is meant in the present invention proteins having a sequence identity, at
5 the amino-acid level, of at least 40%, preferably of at least 50%, more preferably of at least 60% and most preferably of at least 70%.

A particular feature of this method is that essential mycete genes or the corresponding encoded proteins are used
10 as targets for the screening of the substances. The method contemplates that essential *C. albicans* genes as well as functionally similar genes and/or genes homologous in sequence of other pathogenic mycetes or the corresponding encoded proteins may be used as targets.

15 According to one embodiment of the screening method of the invention, mycetes cells are provided, which contain the essential gene used as target, and those cells are incubated with the substance to be tested. By this way, the growth inhibitory effect of this substance with respect to
20 the essential target gene is determined.

The mycetes cells which express the essential target gene to a different degree are used, and these cells are then incubated with the substance to be tested and the growth inhibitory effect of this substance is determined.

25 The method includes the use of two or more mycetes cells, or strains derived therefrom, which differ in that they express the essential target gene to a different degree.

For example, two, three, four, five, ten or more
30 mycetes cells or the corresponding mycetes strains may be comparatively analysed with respect to the growth inhibitory effect of a substance used in a defined concentration. Through such expression degree series, antimycotic substances may be distinguished from cytotoxic
35 or inactive substances.

A particular embodiment of the method includes the use of haploid mycetes cells/strains for the screening, especially haploid *S. cerevisiae* cells/strains.

The method contemplates the integration of the
 5 essential gene selected as a target in a suitable expression vector.

As expression vectors *E. coli*/*S. cerevisiae* shuttle vectors are for example suitable. Especially vectors differing in their copy number per cell may be used.
 10 Therefore, one may use vectors, which are present in the transformed *S. cerevisiae* cells in a high copy number, or one can also use those with a low copy number. One embodiment comprises the use of expression vectors which allow the integration of the target gene in the *S.*
 15 *cerevisiae* genome.

For example the vectors pRS423, pRS424, pRS425, pRS426, pRS313, pRS314, pRS315, pRS316, pRS303, pRS304, pRS305, pRS306 (Sikorki and Hieter, 1989; Christianson et al. 1992) are appropriate as expression vectors.

20 The vectors of the series pRS423-pRS426 are present in a high copy number, about 50-100 copies/cell. On the contrary, the vectors of the series pRS313-pRS316 are present in a low copy number (1-2 copies/cell). When expression vectors from these two series are used, then the
 25 target gene is present as an extrachromosomal copy. Using the vector of the series pRS303-pRS306 allows the integration of the target genes into the genome. Using these three different expression vector types allows a gradual expression of the studied functionally similar
 30 essential gene.

The method includes that the growth inhibitory effect of substances with respect to mycetes cells/strains is comparatively determined using expression vectors differing for instance in the copy number of the vector/cell.

Such cells may express the essential target gene to a different degree and may exhibit a graduated reaction with respect to the substance.

The method includes also, that a target gene
 5 expression of different strength is obtained in different mycetes cells (regulated overexpression) by insertion of the target gene in the expression vector between specific selected *S. cerevisiae* promoters and terminators. *S. cerevisiae* promoters which are constitutively expressed,
 10 but with different strength, are suitable. Examples for such promoters are native promoters of *S. cerevisiae* genes MET25, PGK1, TPI1, TDH3, ADH1, URA3, TRP1, as well as corresponding derivatives therefrom, for example promoter derivatives without specific activator and/or repressor
 15 sequences.

Regulated promoters are also appropriate for the graduated over-expression of the target gene. The native promoters of the GAL1 genes and/or corresponding derivatives thereof, for example promoters, in which different UAS
 20 elements have been eliminated. (GALS, GALL; Mumberg et al. (1994) Nucleic Acids Research 22: 5767-5768) as well as promoters of gluconeogenic genes, for example the promoters FBP1, PCK1, ICL1, or parts thereof, for example their activator- (UAS1 or UAS2) or repressor- (URS) sequences are
 25 used in corresponding non activable and/or non repressible test promoters (Schüller et al. (1992) EMBO J. 11: 107-114) Guarente et al. (1984) Cell 36: 503-511; Niederacher et al. (1992) Curr. Genet. 22: 363-370; Proft et al. (1995) Mol.Gen. Genet. 246: 367-373).

30 In the expression vector terminator for example the terminator sequence of *S. cerevisiae* genes MET25, PGK1, TPI1, TDH3, ADHI, URA3 may be used.

The method includes that by the use of cleverly selected expression vector types and/or the preparation of
 35 suitable expression vectors, eventually using promoters of different strength and differently regulated promoters, a

series of expression vectors may be constructed, all containing the same target gene, but differing in that they express the target gene to a different extent.

The method includes the transformation of the
5 expression vector in haploid wild-type cells of *S. cerevisiae*. The thus obtained *S. cerevisiae* cells/strains are cultivated in liquid medium and incubated in the presence of different concentrations of the tested
10 substance and the effect of this substance on the growth behaviour of the cells/strains expressing the target gene to a different degree is comparatively analysed. The method also includes that haploid *S. cerevisiae* cells/strains, transformed using the respective expression vector type without target gene, are used as a reference.

15 The method includes that the screening of the substances can be carried out in different media using regulated promoters, especially GAL1 promoter and its derivatives (GALS and GALL), since the expression degree may be largely influenced by the choice of the respective
20 medium. Thus, the expression degree of the GAL1 promoter decreases in the following fashion: 2 % galactose > 1 % galactose + 1 % glucose > 2 % glycerine > 2 % glucose.

The effect of the substances inhibiting the growth of wild-type cells of *S. cerevisiae*, may be partially or
25 totally compensated by the overexpression of the functionally similar gene of another mycete species.

According to one embodiment, the method for screening antimycotic substances is carried out in vitro by contact
30 of an essential or functionally similar gene or the corresponding encoded protein with the substance to be tested and determination of the effect of the substance on the target. Any in vitro test appropriate for determining the interaction of two molecules, such as a hybridization test or a functional test, can be used (e. g. enzymatic
35 tests which are described in details in Bergmeyer H. U., Methods of Enzymatic Analysis, VCH Publishers). If the

screening is carried out using the encoded protein as the target, then the corresponding essential gene is inserted by any suitable method known in the art, such as PCR amplification using a set of primers containing appropriate restriction sites, (Current Protocol in Molecular Biology, 5 John Wiley and Sons, Inc) into an expression system, such as *E. coli*, Baculovirus, or yeast, and the expressed protein is then completely or partially purified by a method known in the art. Any purification method 10 appropriate for the purification of expressed proteins, such as affinity chromatography can be used. If the target protein function is known, a functional test can then be carried out in which the effect of the antimycotic substance on the protein function is determined. If the 15 protein function is unknown, substances which can interact with the target protein, e. g. which bind to the encoded protein, can be tested. In such a case a test such as protection of the target protein from enzymatic digestion by appropriate enzymes can be used.

20 According to one specific embodiment, the method for screening antimycotic substances corresponds to an enzymatic assay wherein the activity of dihydropteroate synthase (DHPS) and/or 7,8-dihydro-6-hydroxymethylpterin-pyrophosphokinase (HPPK) is determined; the enzymatic essay 25 can be such as disclosed in "Bergmeyer H. U., Methods in Enzymatic analysis, VCH Publishers".

Dihydropteroate synthetase (DHPS) catalyses the condensation of 6-hydroxymethyl-7,8-dihydropterin pyrophosphate to para-aminobenzoic acid to form 7,8- 30 dihydropteroate which corresponds to the second step in the three-step pathway leading from 6-hydroxymethyl-7,8-dihydropterin to 7,8-dihydrofolate. 7,8-dihydro-6-hydroxymethylpterin-pyrophosphokinase (HPPK) catalyzes the attachment of pyrophosphate to 6-hydroxymethyl-7,8- 35 dihydropterin to form 6-hydroxymethyl-7,8-dihydropterin pyrophosphate which corresponds to the first step in a

three-step pathway leading to 7,8-dihydrofolate. All organisms require reduced folate cofactors for the synthesis of a variety of metabolites. Most microorganisms must synthesize folate de novo because they lack the active
5 transport system of higher vertebrate cells which allows these organisms to use dietary folates. Enzymes involved in folate biosynthesis are therefore targets for a variety of antimicrobial agents. Consequently, these enzyme activities are essential to the microorganisms, and are absent in man.

10 The method also includes the identification of genes which are functionally similar and/or homologous in sequence to essential *C. albicans* genes from humans, animals or plants. The corresponding human, animal or plant genes may optionally be used as target genes in the method
15 in order to test if antimycotic substances exhibit an effect on these target genes.

A particular advantage of the method is that in this way substances may be identified which efficiently inhibit mycetes growth and also the influence of these substances
20 on corresponding functionally similar genes and/or genes homologous in sequence to essential *C. albicans* genes from human, animal or plants may be determined.

The method includes also the possibility to check the existence of functionally similar genes and/or human,
25 animal or plant genes homologous in sequence to the corresponding essential mycete genes, for example by checking homology of the identified essential mycete genes or parts thereof with human, animal or plant sequence genes available in data banks. In this way, it is possible to
30 select at an early stage from the identified essential mycete genes, depending on the aim, those for which no functionally similar gene and/or no human gene homologous in sequence exist, for example.

Thereby, the method offers a plurality of
35 possibilities to identify selectively substances with

antimycotic effects, with no harmful effect on human beings.

For example, it is possible to identify substances usable for the preparation of drugs for the treatment of mycosis or prophylaxis in immunodepression states. These substances may be used for example for the manufacture of drugs usable for the treatment of mycotic infections, which occur during diseases like AIDS or Diabetes. Substances which may be used for the fabrication of fungicides, especially of fungicides which are harmless for humans and animals, can also be identified.

Furthermore, the method offers the possibility to identify antimycotic substances, which selectively inhibit growth of specific mycete species only.

The screening method is particularly advantageous inasmuch as it is sufficient to know whether the genes are essential, one does not need any additional information regarding the function of the essential genes or the function of the encoded proteins. In addition, it is particularly advantageous for the identification of functionally similar genes to essential *S. cerevisiae* gene, in other mycetes where the DNA sequence is not available for many of these genes.

According to another aspect the invention provides an antibody directed against the protein encoded by the CaOR110, CaMR212, CaNL256, CaBR102, CaIR012, CaDR325 or CaJL039, gene or a polypeptidic fragment thereof. The term "antibody" encompasses monoclonal and polyclonal antibodies. Said antibodies can be prepared by method well known in the art such as those disclosed in "Antibodies, a laboratory manual", Ed. Harbow and David Lane. Cold Spring Harbor Laboratory Eds., 1988.

According to another aspect the present invention provides a kit for the diagnosis of fungal infections comprising a gene selected from the group consisting of CaOR110, CaMR212, CaNL256, CaBR102, CaIR012, CaDR325 and

CaJL039, a functionally similar gene thereof, a functional fragment thereof, the corresponding encoded protein, a functional polypeptide fragment thereof or an antibody directed against the protein encoded by CaOR110, CaMR212, CaNL256, CaBR102, CaIR012, CaDR325 or CaJL039 gene or by a functionally similar gene, or a polypeptidic fragment thereof. Such kits can be prepared using any suitable method well known in the art.

10 Examples

Example 1: CaNL256

The Internet site of Stanford (<http://candida.stanford.edu/>) gives access to preliminary sequences of the genome of *C. albicans*. One of these sequences has homology with the YNL256 gene of *S. cerevisiae*. Two oligonucleotides were selected in this sequence (5'-ATTCATCCCATCAGTGCAGAAAG-3' (SEQ ID NO:27) and 5'-ATTGACCAATAGCTCTAATTAATG-3') (SEQ ID NO:28) in order to amplify the corresponding fragment of *C. albicans*. After cloning, we obtained a sequence of 399 bp close to the expected sequence (SEQ ID NO: 1). The deduced protein was compared with the one of YNL256, evidencing 53% similarity and 43% identity (fig. 1). This fragment of 399 bp of *C. albicans* was used as a probe for screening a genomic library of *C. albicans*. The latter was prepared by partial digestion of genomic DNA of *C. albicans* by Sau3AI and cloning into the YEP24 vector at the BamHI site. The clones of the library were then spread at a density of 2000 clones per dish. Each dish was covered by a nitrocellulose filter which was then successively treated with: NaOH, 0.5M, 5 minutes; Tris, 1M, pH 7.7, 5 minutes; Tris, 0.5M, pH 7.7, NaCl, 1.25M, 5 minutes. After drying, the filters were kept for 2 hours at 80°C. Prehybridization and hybridization were carried out in a buffer of 40% formamide, 5xSSC, 20 mM Tris pH 7.7 1xDenhardt 0.1% SDS. The probe was labeled with ³²P with the RediPrime kit and dCTP from Amersham

UK. Hybridization took place over 17 hours at 42°C. The filters were then washed in 1x SSC, 0.1% SDS, three times for 5 minutes at room temperature and then twice for 30 minutes at 60°C, and were then submitted to autoradiography overnight. The colonies corresponding to the spots obtained were reisolated by re-spreading at low density followed by further hybridization. Three clones were thus obtained (out of 40,000), which were sequenced on an ABI 377 apparatus. The sequences were compiled using the ABI software and then analysed using the GCG software package. One of these three clones turned out to contain the complete coding sequence corresponding to the probe used; this gene was called CaNL256, whose sequence is represented in SEQ ID NO: 2. CaNL256 has 52% of nucleotides identical to YNL256 of *S. cerevisiae*. The coding region is shorter at the N-terminus. For translation to amino acids, account was taken of the fact that, in *C. albicans*, the CTG codon is translated to Serine (there are 3 CTG codons in CaNL256). The deduced protein (SEQ ID NO: 3) had 40% amino acids identical with YNL256 of *S. cerevisiae* and 41% with FAS (folic acid synthase) of *Pneumocystis carinii*. Investigation into the databases using the Blast software showed homology of two parts of the CaNL256 protein with, respectively, the bacterial enzymes Dihydropteroate Synthase (EC 2.5.1.15) (DHPS) of *Haemophilus influenzae*, *Staphylococcus haemolyticus*, *Neisseria meningitidis*, *Streptococcus pneumoniae*, *Bacillus subtilis*, *Clostridium acetobutylicum*, *Escherichia coli*, *Mycobacterium leprae* (P value less than e^{-28}) and 7,8-dihydro-6-hydroxymethylpterin-pyrophosphokinase (EC 2.7.6.3) (HPPK) of *Bacillus subtilis*, *Escherichia coli*, *Haemophilus influenzae*, *Streptococcus pneumoniae* (P value less than e^{-20}). The units characteristic of DHPS and HPPK are also found in CaNL256.

Example 2: CaBR102

35 The Internet site of Stanford

(<http://candida.stanford.edu/>) give access to preliminary sequences of the genome of *C. albicans*. One of these sequences has homology with the YBR102 gene of *S. cerevisiae*. Two oligonucleotides were selected in this

5 sequence (5'-AGTATTCAATTGGGTATTCC-3' (SEQ ID NO:29) and 5'-CCGGCATCATCAGTAACTCC-3') (SEQ ID NO:30) in order to amplified the corresponding fragment of *C. albicans*. After cloning, we obtained a sequence of 647 bp (SEQ ID NO: 4). The deduced protein was compared with the one of YNL102,

10 evidencing 35% similarity and 26% identity (fig. 2). This fragment was amplified using Pfu polymerase (Stratagene). The PCR product was purified (High Pure PCR Product Purification Kit, Boehringer Mannheim) and used as a probe for screening a *C. albicans* genomic DNA library. The latter

15 was prepared by partial digestion of *C. albicans* genomic DNA with *Sau*III A and cloning into the YEP-24TRP1 vector at the *Bam*HI restriction site. 40,000 clones of the library were then spread at a density of 2000 clones per dish. Each dish was covered by a nitrocellulose filter (Membrane

20 Hybond N', Amersham) which was then successively treated with : 1.5 M NaCl/0.5 M NaOH, 5 minutes; 1.5 M NaCl/0.5 M Tris-HCl pH 7.2/1 mM EDTA, 3 minutes, twice; DNA was crosslinked to the filters (Amersham Life Science, ultra violet crosslinker). The probe (100 ng) was labelled with

25 ³²p using the RediPrime kit and dCTP (Amersham Life Science). The filters were hybridized in a buffer containing 30% formamide, 5 x SSC, 5% Denhart's solution, 1% SDS, 100 yg/ml salmon sperm DNA and a probe concentration of 10⁶ cpm/ml at 42°C for 16 h. The membranes

30 were then washed three times at room temperature in 2 x SSC/0.1% SDS for 5 minutes each and three times in 1 x SSC/0.1% SDS at 60°C for 20 minutes each. the filters were then exposed overnight to an X-ray film. The colonies corresponding to the positives clones were isolated and

35 screened a second time by the same procedure. Two positives clones were finally obtained, which were sequenced on an

ABI377 apparatus. the sequences were compiled using ABI software and then analysed using the GCG software package. The nucleotide sequences of these two clones were identical and contained the complete coding sequence corresponding to the probe used, this gene was called CaBR102, whose sequence is represented in SEQ ID No: 5 and 6. The translation of this nucleotide sequence was examined, account was taken of the fact that in *C. albicans* the CTG codon is translated to serine (there are 3 CTG codons in CaBR102). The deduced protein has 24% identity to *S. cerevisiae* gene YBR102.

Example 3: CaIR012

Chromosomal DNA from the *C. albicans* strain Caf2-1 was isolated using Yeast Cell Lysis prep Kit and Genome DNA Kit from BI0101. A 343 bp fragment from *C. albicans* genomic DNA (SEQ ID NO: 7) was amplified with the oligonucleotide primers CaYIR012-5' (5'-GACGTCGTAGACGATACTCAAGAAG-3') (SEQ ID NO:31) and CaYIR012-3' (5'-CTGCAGTAAACCCTCCAGATATAACAG-3') (SEQ ID NO:32) by PowerScript DNA polymerase (PAN Systems GmbH) using the hot start technique. The PCR product was purified from the agarose gel and labeled with fluorescein (Gene image random prime labelling module, Amersham Life Science) according to the manufacturer's instructions. Plasmid DNA from *E. coli* was isolated using Qiagen columns as recommended by the manufacturer. Screening the kZAPII *C. albicans* cDNA library was performed following the manufacturer's instructions (Stratagene Ltd.). Nylon filters (Schleicher&Schuell) were lifted from LB-plates (150 mm) with 15000 pfu/plate, denatured 5 min in 1.5M NaCl, 0.5M NaOH, neutralized 3 min in 1.5M NaCl, 0.5M Tris-HCl pH8.0, washed 3 min in 0.2 M Tris-HCl pH 7.5, 2xSSC and DNA was crosslinked to the filters (Stratagene W crosslinker). The filters were prehybridized 4 h at 60 °C and hybridized with the fluorescein-labeled DNA probe overnight at 60 °C. Detection was performed with Anti-fluorescein AP conjugate (Signal amplification module for

the FluorImager, Amersham LIFE SCIENCE) and analysed after 20 h with a Fluorimager (Storm 860, Molecular Dynamics). Positive plaques were picked and incubated with 0.5 ml SM-buffer (100mM NaCl, 8 mM MgSO₄, 50 mM Tris-HCl pH7.5, 0.01% gelatin). The selected clones were diluted, titered with host cells XL1- Blue and screened and purified a second time by the same procedure. Finally, the pBluescript SK (-) phagemid containing the DNA insert of interest was rescued by the ExAssist Helper Phage system according to the Stratagene protocol. From a total of 75000 screened plaques, 3 positive clones were identified. pBluescript SK (-) phagemid DNA was isolated, sequenced with T3 and T7 primers and the sequences were extended with custom-synthesized oligonucleotide primers. Nucleotide sequence analyses were performed with the Gene Data software package (Gene Data AG, Basel Switzerland). Similarity searches with the Swissprot database were conducted with the BLAST program (Gish, Warren and David J. States (1993). Identification of protein coding regions by database similarity search. Nat. Genet. 3: 266-72.). One of these three clones turned out to contain the complete coding sequence corresponding to the probe used; this gene was called CaIR1012, whose sequence is represented in SEQ ID NO: 8 and 9.

Example 4: CaJL039

The CaJL039 sequence is depicted in SEQ ID No 10 and 11.

The CaJL039 gene was cloned based on gene fragment data issued from the public Stanford Candida albicans sequencing database.

(a) A fragment that showed homology to *Saccharomyces cerevisiae* YJL039c was identified, the sequence of which is given in SEQ ID No 12.

Using the procedure disclosed in example 3 with the oligonucleotide primer pair (Ca039s: TAG CTC AAC CTA CCA CCA ATC (SEQ ID NO:33)/Ca039r: ATC ACA AGA CTG TCA ATG TAA

AT) (SEQ ID NO:34), a short PCR fragment (234 base pairs long) was amplified for screening a *Candida albicans* cDNA lambda ZAP II library (gift of Alistair Brown, Aberdeen).

Three positive clones of the 3 coding region were
5 obtained. [(#)21t7, 11t3, 21t3).

(b) 3-and 5-extension of the internal fragment using the primer walking method

The Sanglard genomic *Candida* DNA library with the YEp24 vector backbone was used for further amplification of
10 3-and 5-coding sequences. Amplification was carried out by using the following vector-specific oligonucleotide primers and CaJL039 fragment-specific primers:

Cggaattcctatcgactacgcatcatgg (SEQ ID NO:35): YEp24for (vector specific)

15 Gcgaattccgatataggcgccagcaac (SEQ ID NO:36): YEp24ba (vector specific)

Caattgctttgactcgggtgttattaagt (SEQ ID NO:37): Ca039-51 (CaJL039: 5fishing)

Tcttggcacaacttgataagaatctgt (SEQ ID NO:38): Cl039-52

20 Taggtgtacgcgaaagccaagtagaac (SEQ ID NO:39): Ca039-53 ()

Ttgttaatcgtacacctaaggtgttgac (SEQ ID NO:40): Ca039-31 (CaJL039: 3fishing)

Ttgcagattgatgctagcaatgtatttg (SEQ ID NO:41): Ca039-32

Using the technique of primer walking, the complete
25 5'-sequence could be amplified (clone 14b-1-1 and clone 17b-3-4).

The missing 3-sequence was available from GTC PathoGenome Release 5.0, contig #2830.

An interacting protein (C82, component for RNA
30 polymerase III in yeast) has been identified.

Example 5: CaOR110

5.1. CaOR110

The CaOR110 sequence is depicted in SEQ ID No 13 and
14.

35 CaOR110 was cloned based on gene fragment data issued from the public Stanford *Candida albicans* sequencing database.

(a) A small ScOR110-homologous fragment was used in a hybridization experiment to identify CaOR110 clones in a Candida Albicans lambda ZAPII cDNA library (from Alistair Brown). Alignment of Candida Albicans CaOR110 sequence with the fragment used for hybridization is given in figure 3. The homologous fragment sequence is given in SEQ ID No.25.

(b) 3'-and 5'-Extension of the internal fragment: The Sanglard genomic Candida DNA library (received from RMV) in the YEP24 vector backbone was used for the amplification of 3'-and 5'-coding and non-coding sequences. This amplification was done by using the vector- specific oligos (directional towards the insert) and CaOR110 fragment-specific oligos (directional towards the vector flanking sequences) described below:

cggaattcctatcgactacgcatcatgg (SEQ ID NO:35) : YEP24for
gcgaattccgatataggcgccagcaac (SEQ ID NO:36) : YEP24ba
cgggatccggtaaccaattggatctataaccgtg (SEQ ID NO:42) :
110-ba-150
gcggatcctggtgcccttggtggtgaatg (SEQ ID NO:43) :
CaYOR110A
gcggatccctcacaatatgacgattgaaact (SEQ ID NO:44) :
CaYOR110B
ggcgtcgactcaggcgccagttttacgtacttcaaattcatc (SEQ ID
NO:45) : CaYOR110C
tgtgaattcttgacacaggggtga (SEQ ID NO:46) : CaYOR110D
caaaccttcagcacaactcca (SEQ ID NO:47) : CaYOR110E

The finally assembled sequence that included also 3'-and 5'-non-coding sequences was verified by sequencing.

The coding region was subcloned into the p414RSGALL-vector. The map is depicted in Fig. 4.

The homologous yeast ORF (YOR110w) has been described as the transcription factor subunit TFC7 interacting with TFC1 in the TFIIIC polymerase complex (Manaud et al., 1998, Mol. Cell. Biol. 18; 3191-3200).

5.2. CaOR110 splice variant

For CaOR110, an additional splice variant was identified. The clones for the splice variant of CaOR110 were obtained from a *Candida albicans* cDNA library.

The sequence is depicted in SEQ ID No. 15 and 16.

5 The splice variant uses the donor site "gtacgt" at position 907 of the original CaOR110 sequence. Acceptor site is at 1047. The map is disclosed in Fig. 5.

The alignment of the original CaOR110 and the splice variant is given in fig. 6.

10 Example 6: CaMR212

The CaMR212 sequence is depicted in SEQ ID No. 17 and 18

(a) CaMR212 was cloned based on gene fragment data from the public Stanford *Candida albicans* sequencing database.

The sequence of a fragment showing homology (Blast search) to the *Saccharomyces cerevisiae* gene YMR212c is given in SEQ ID 19.

Based on these data, the following oligos were designed that allow amplification of this fragment (490 bp-fragment) from genomic *Candida albicans* DNA. Oligos:

CaYMR212for: 5'-cacctgtgaacaacccaccatc-3 (SEQ ID NO:48)

CaYMR212back: 5'-gaatatcctttttaactcaagag-3' (SEQ ID NO: 49)

(b) 3'-and 5'-extension of this internal fragment from CaMR212

For this purpose, genomic *Candida* DNA libraries from Dominique Sanglard (received from RMV) were used. The YEp24 backbone of this library was used to amplify the 3'-and 5'-coding and non coding sequences with PCR. This was done by using oligos specific for the CaMR212 490 bp-fragment (directional towards the vector flanking regions) and vector-specific oligos (directional towards the insert).

35 Oligos:

YEP24for (vector specific):

5'-cggaattcctatcgactacgcgatcatgg (SEQ ID NO: 35) YEP24ba (vector specific):

5'-gcgaattccgatataggcgccagcaac (SEQ ID NO: 36)

5 Primer YEp24for and CaMR212for gave a 500 bp fragment, encoding 5'-UTR and the 5'coding region from CaMR212.

Using primer YEp24 for and CaMR212back a 1400 bp CaMR212-fragment was amplified. Using the sequence of this 1400 bp-fragment the following new primers, specific for
10 this fragment were designed.

Oligos:

Ca212-1: 5'-gctttcccagcaggataaacattg (SEQ ID NO:50)

Ca212-2: 5'-tgagttataatgcagctgttgg (SEQ ID NO:51)

15 Ca212-3: 5'-catctcgtgtgaacatgattgg (SEQ ID NO:52)

Primers YEP24 for and Ca212-3 gave a 1600 bp fragment, coding for the 3'-coding region and the 3'UTS region.

With the 3 PCR fragments the 2900 bp sequence (including coding and 3'and 5'-non-coding sequences) was
20 assembled. With the following new primers the coding sequences was amplified from genomic DNA and cloned into p413GALL-vector.

Oligos for amplifying coding region:

Ca212for: 5'-agtttcttcaacttccagatccaag (SEQ ID NO: 53)

25 Ca212back: 5'-gtatatttgcaactgtctctctctc (SEQ ID NO: 54)

The yeast homolog YMR212c plays a role in cell wall function because the knockout can be rescued in 1M sorbitol. In addition, YMR212c under GAL-promoter
30 regulation shows an increased sensitivity versus Congo Red and Calcofluor White. YMR212c is an integral membrane protein and localizes to the plasma membrane (demonstrated by microscope analysis of YMR212-GFP fusion proteins and by biochemical analysis of YMR212-GST fusion proteins).

35 Example 7: CaDR325

The CaDR325 sequence is given at SEQ ID 20 and 21.

CaDR325 was cloned based on gene fragment data from the public Stanford *Candida albicans* sequencing database.

(a) 3 fragments that showed homology to *Saccharomyces cerevisiae* YDR325 were identified, the sequences of which are disclosed in SEQ ID 22, 23 and 24.

Based on these data, the following oligos were designed that allowed the verification of the database sequences and the amplification of an approx. 2200 bp internal CaDR325 fragment from genomic DNA:

10 cgagcatctacttgttcaaccac: hybCaYDR325ba Oligo(SEQ ID NO: 55)

gaatctctggctcgctc: 325-juls Oligo (SEQ ID NO:56)

gaccgagatacagagaat: 325-julr Oligo(SEQ ID NO:57)

ggttaaattgatcgtgatgaat: Ca325r Oligo (SEQ ID NO:58)

15 caacctcactgacaaatactt: Ca325s Oligo (SEQ ID NO:59)

The finally subcloned 2200 bp internal fragment was amplified by the combination hybCaYDR325ba + 325-julr oligos.

(c) 3'-and 5'-Extension of the internal fragment: The Sanglard genomic *Candida* DNA library (received from RMV) in the YEP24 vector backbone was used for the amplification of 3'-and 5'-coding and non-coding sequences. This was done by using the following vector- specific oligos (directional towards the insert) and CaDR325 2200 bp fragment-specific oligos (directional towards the vector flanking sequences):

25 cggaattcctatcgactacgcatcatgg (SEQ ID NO:35) :
YEP24for (vector specific)

gcgaattccgatataggcgccagcaac (SEQ ID NO:36) : YEP24ba
(vector specific)

30 acgcttccaatgtattattctcg (SEQ ID NO:60) : Oligo 1-10-A
back

ggatgccaatctccctga(SEQ ID NO: 61) : Oligo 1-10-B for

catccagaagatataacggct (SEQ ID NO: 62) : Oligo 1-10-C
for

35 tgcataatctactcagcgaca (SEQ ID NO:63) : Oligo 1-10-D
back

gtggttgaacaagtagatgctcg (SEQ ID NO:64) : Oligo 1-10-E
for
gcgcttgaaaccactagtgattg (SEQ ID NO:65) : Ca325Klon
2_Fo

5 caattcactagtggtttcaagcgc (SEQ ID NO: 66) :
Ca325Klon3Ba

The finally assembled 4700 bp sequence that included
also 3'-and 5'-non-coding sequences were verified by
sequencing. The coding region was subcloned into the
10 p413RSGALL-vector.

The map is disclosed in fig. 7.

Sequences numbers are identified in field 130 of the
sequence listing.